

UNITED STATES DISTRICT COURT
DISTRICT OF NEW JERSEY

<p>IN RE: VALSARTAN, LOSARTAN, AND IRBESARTAN PRODUCTS LIABILITY LITIGATION</p>	<p>MDL No. 2875</p>
<p>THIS DOCUMENT RELATES TO ALL CASES</p>	<p>HON. ROBERT B. KUGLER</p>

NOTICE TO TAKE VIDEOTAPED ORAL DEPOSITION

TO: **Seth A. Goldberg, Esq.,
DUANE MORRIS LLP
30 South 17th Street
Philadelphia, Pennsylvania 19103**
Attorneys for Defendants Zhejiang Huahai Pharmaceutical Co, Ltd., Huahai U.S., Inc., Princeton Pharmaceutical Inc., and Solco Healthcare US, LLC (hereinafter "Defendants").

Please take notice that pursuant to Federal Rule of Civil Procedure 30, and other applicable Rules, including the Local Civil Rules, and the applicable Orders of the Court, Plaintiffs, by and through their counsel, will take the videotaped deposition of Lihong (Linda) Lin, Director of Regulatory Affairs at Zhejiang Huahai Pharmaceutical Co., Ltd., on May 4 through 7, 2021, at 7:00 a.m. Hong Kong time (May 3 through 6, 2021, at 7:00 p.m. eastern time), and continuing until completion, at Duane Morris LLP, 30 South 17th Street, Philadelphia, Pennsylvania 19103, via Zoom, in accordance with the Fact Witness Deposition Protocol, Case Management Order #20, filed November 17, 2020 (Document 632). The deposition shall first address the Federal Rule of Civil Procedure 30(b)(6) topics listed on Exhibit A attached hereto, followed by deposition of the witness in her individual capacity. The witness shall produce the documents requested at Exhibit B, attached hereto, at least five days in advance of the deposition.

Pursuant to the meet and confer between the parties, a translator will be provided.

TAKING ATTORNEYS FOR PLAINTIFFS:

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The videotaped deposition will be taken before a person authorized by law to administer oaths, pursuant to Rule 28 of the Federal Rules of Civil Procedure.

April 15, 2021

PLAINTIFFS' CO-LEAD COUNSEL

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EXHIBIT A

30(B)(6) TOPICS

On behalf of Zhejiang Huahai Pharmaceutical Co., Ltd:

Communications with Regulatory Agencies

38. The communications with any regulatory authority, including but not limited to the FDA, with regard to the modifications with regard to the use of solvents, and the Tetrazole ring formation step, in the manufacturing process for ZHP's valsartan API.

39. The communications with any regulatory authority, including but not limited to the FDA, with regard to the modifications with regard to the use of solvents, and the Tetrazole ring formation step, in the manufacturing process for ZHP's finished dose.

40. ZHP's disclosures to regulatory authorities, including the FDA, with regard to the actual or potential contamination of ZHP's valsartan API with nitrosamines including NDMA and NDEA.

41. ZHP's filings with regulatory authorities, including the FDA, regarding manufacturing process changes for ZHP's Valsartan API Drug Master Filings.

EXHIBIT B

DOCUMENT REQUESTS

1. The most recent resume/Curriculum Vitae and LinkedIn profile for Lihong (Linda) Lin.
2. The complete production of Lihong (Linda) Lin's relevant custodial documents, including those maintained on personal computers or electronic devices, to the extent not produced prior.

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CERTIFICATE OF SERVICE

I hereby certify that on April 15, 2021, I electronically filed the foregoing document with the Clerk of the court using CM/ECF system which will send notification of such filing to the CM/ECF participants registered to receive service in this MDL.

PLAINTIFFS' CO-LEAD COUNSEL

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